=> d ibib abs hitstr 1-2 THE ESTIMATED COST FOR THIS REQUEST IS 11.28 U.S. DOLLARS DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y) /N:v

ANSWER 1 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2008:1169030 CAPLUS

DOCUMENT NUMBER: 149:493846

TITLE: 5-N-Methylated Ouindoline Derivatives as Telomeric G-Ouadruplex Stabilizing Ligands: Effects of 5-N Positive Charge on Quadruplex Binding Affinity and

Cell Proliferation

AUTHOR(S): Lu, Yu-Jing; Ou, Tian-Miao; Tan, Jia-Heng; Hou,

Jin-Qiang; Shao, Wei-Yan; Peng, Dan; Sun, Ning; Wang, Xiao-Dong; Wu, Wei-Bin; Bu, Xian-Zhang; Huang,

Zhi-Shu; Ma, Dik-Lung; Wong, Kwok-Yin; Gu, Lian-Quan CORPORATE SOURCE: School of Pharmaceutical Sciences, Sun Yat-sen University, Guangzhou, 510080, Peop. Rep. China Journal of Medicinal Chemistry (2008), 51(20),

SOURCE: 6381-6392

CODEN: JMCMAR; ISSN: 0022-2623 PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal

LANGUAGE: English OTHER SOURCE(S): CASREACT 149:493846

AB A series of 5-N-Me quindoline (cryptolepine) derivs. as telomeric quadruplex ligands was synthesized and evaluated. The designed ligands possess a pos. charge at the 5-N position of the aromatic quindoline scaffold. The quadruplex binding of these compds. was evaluated by CD (CD) spectroscopy, fluorescence resonance energy transfer (FRET) melting assay, polymerase chain reaction (PCR) stop assay, NMR (NMR), and mol. modeling studies. Introduction of a pos. charge not only significantly improved the binding ability but also induced the selectivity toward antiparallel quadruplex, whereas the nonmethylated derivs. tended to stabilize hybrid-type quadruplexes. NMR and mol. modeling studies revealed that the ligands stacked on the external G-guartets and the pos. charged 5-N atom could contribute to the stabilizing ability. Long-term exposure of human cancer cells to I showed a remarkable cessation in population growth and cellular senescence phenotype and accompanied by a shortening of the telomere length.

TT 1072837-74-1P 1072837-78-5P 1072837-81-0P 1072837-83-2P 1072837-99-0P 1072838-00-6P 1072838-01-7P 1072838-02-8P 1072838-03-9P 1072838-04-0P 1072838-05-1P 1072838-06-2P RL: PAC (Pharmacological activity); PRP (Properties); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation) (preparation and evaluation of cryptolepine derivs.)

RN 1072837-74-1 CAPLUS

CN 1,2-Ethanediamine, N1,N1-dimethyl-N2-(5-methyl-5H-quindolin-11-yl)-, hydriodide (1:1) (CA INDEX NAME)



Me2N-CH2-CH2-NH

HI

1072837-78-5 CAPLUS

1,3-Propanediamine, N1,N1-dimethyl-N3-(5-methyl-5H-quindolin-11-yl)-, CN hydriodide (1:1) (CA INDEX NAME)



Me2N- (CH2)3-NH

HI

RN 1072837-81-0 CAPLUS

CN 1,2-Ethanediamine, N1,N1-diethy1-N2-(5-methy1-5H-quindolin-11-y1)-, hydriodide (1:1) (CA INDEX NAME)

Et2N-CH2-CH2-NH

HI

- RN 1072837-83-2 CAPLUS
- CN 1,3-Propanediamine, N1,N1-diethyl-N3-(5-methyl-5H-quindolin-11-yl)-, hydriodide (1:1) (CA INDEX NAME)



Et2N- (CH2)3-NH

• HI

- RN 1072837-99-0 CAPLUS
- CN 1,2-Ethanediamine, N2-(7-fluoro-5-methyl-5H-quindolin-11-yl)-N1,N1-dimethyl-, hydriodide (1:1) (CA INDEX NAME)

Me2N-CH2-CH2-NH

HI

- RN 1072838-00-6 CAPLUS
- CN 1,3-Propanediamine, N3-(7-fluoro-5-methyl-5H-quindolin-11-yl)-N1,N1dimethyl-, hydriodide (1:1) (CA INDEX NAME)

Me2N- (CH2)3-NH

HI

- RN 1072838-01-7 CAPLUS
- CN 1,2-Ethanediamine, N1,N1-diethyl-N2-(7-fluoro-5-methyl-5H-quindolin-11-yl)-, hydriodide (1:1) (CA INDEX NAME)

Et2N-CH2-CH2-NH

HI

RN 1072838-02-8 CAPLUS

CN 1,3-Propanediamine, N1,N1-diethyl-N3-(7-fluoro-5-methyl-5H-quindolin-11-yl)-, hydriodide (1:1) (CA INDEX NAME)

Et2N- (CH2)3-NH

• HI

RN 1072838-03-9 CAPLUS

CN 1,2-Ethanediamine, N2-(7,9-difluoro-5-methyl-5H-quindolin-11-yl)-N1,N1dimethyl-, hydriodide (1:1) (CA INDEX NAME)

Me2N-CH2-CH2-NH

HI

RN 1072838-04-0 CAPLUS

CN 1,3-Propanediamine, N3-(7,9-difluoro-5-methyl-5H-quindolin-11-yl)-N1,N1-dimethyl-, hydriodide (1:1) (CA INDEX NAME)

## HI

RN 1072838-05-1 CAPLUS

CN 1,2-Ethanediamine, N2-(7,9-difluoro-5-methyl-5H-quindolin-11-yl)-N1,N1-diethyl-, hydriodide (1:1) (CA INDEX NAME)

Et2N-CH2-CH2-NH F

## HT

RN 1072838-06-2 CAPLUS

CN 1,3-Propanediamine, N3-(7,9-difluoro-5-methyl-5H-quindolin-11-yl)-N1,N1-diethyl-, hydriodide (1:1) (CA INDEX NAME)

• HI

REFERENCE COUNT: 67

67 THERE ARE 67 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2005:523449 CAPLUS

DOCUMENT NUMBER: 143:59844

TITLE: A preparation of antitumor quinoline derivatives

INVENTOR(S): Aymami Bofarull, Juan; Coll Capella, Miquel; Llebaria

Soldevila, Amadeo; Navarro Munoz, Isabel

PATENT ASSIGNEE(S): Crystax Pharmaceuticals S.L., Spain; Consejo Superior de Investigaciones Cientificas; Universitat

Politecnica de Catalunya

PCT Int. Appl., 39 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

LANGUAGE: Enc FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

GI

PATENT NO.			IND	DATE		APPLICATION NO.								
			A1 20050616			WO 2004-EP13106								
W:	AE, AG,	AL, A	M, AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ,	CA,	CH,
	CN, CO,	CR, C	U, CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,
	GE, GH,	GM, H	R, HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KZ,	LC,
	LK, LR,	LS, L	T, LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NA,	NI,
	NO, NZ,	OM, P	G, PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,
	TJ, TM,	TN, T	R, TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW
RW:	BW, GH,	GM, K	E, LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,
	AZ, BY,	KG, K	Z, MD,	RU,	TJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,
	EE, ES,	FI, F	R, GB,	GR,	HU,	IE,	IS,	IT,	LU,	MC,	NL,	PL,	PT,	RO,
	SE, SI,	SK, T	R, BF,	BJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,
	NE, SN,	TD, T	G											
EP 1687304			A1 20060809			EP 2004-797988					20041118			
R:	AT, BE,	CH, D	E, DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,
	IE, SI,	FI, R	O, CY,	TR,	BG,	CZ,	EE,	HU,	PL,	SK,	IS			
JP 2007511573			T	20070510			JP 2006-540327				20041118			
US 20070105784			A1 20070510			US 2006-580140					20060519			
PRIORITY APPLN. INFO.:							ES 2003-2821					A 20031120		
							WO 2	004-1	EP13	106	1	7 2	0041	118
OTHER SOURCE	M	MARPAT 143:59844												

## \* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

AB The invention relates to a preparation of antitumor quinoline derive. of formula G1-L-G2 [wherein: G1 is quinoline derivative, L is a single bond or a linking radical; G2 is H, quinoline derivative, or C-9 radical of acridine, etc.]. The invention compds. are intercalators, compds. that bind between DNA base pairs. The in vitro cytotoxicity of the compds. was evaluated by colorimetric assays with tetrazole salts on Jurkat clone E6-1 and on GCC-4, human leukemia and carcinoma cell lines. For instance, indoloquinoline derivative I (Jurkat clone E6-1, IC50 = 1.42 µM) was prepared via amidation of OH-indolo[3,2-b]-1-carboxylic acid by carbamic acid derivative II, decarboxylation of the obtained amide, and subsequent amidation of 9-acridinecarboxylic acid by the obtained amine III (yields: 1st amidation - 70%, decarboxylation - 92%, 2nd amidation - 50%).

IT 854190-47-9P 854190-53-7P 854190-56-0P

854190-70-8P 854190-76-4P 854190-81-1P

854190-92-4P 854190-96-8P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of antitumor quinoline derivs.)

- RN 854190-47-9 CAPLUS
- CN 5H-Quindoline-11-carboxamide, N,N'-[(methylimino)di-3,1-propanediyl]bis[5-methyl- (9CI) (CA INDEX NAME)

- RN 854190-53-7 CAPLUS
- CN 1,2-Ethanediamine, N1,N1-dimethyl-N2-(5-methyl-5H-quindolin-11-yl)- (CA INDEX NAME)



Me2N-CH2-CH2-NH

- RN 854190-56-0 CAPLUS
- CN 1,3-Propanediamine, N-methyl-N'-(5-methyl-5H-quindolin-11-yl)-N-[3-[(5-methyl-5H-quindolin-11-yl)amino]propyl]- (9CI) (CA INDEX NAME)

- RN 854190-70-8 CAPLUS
- CN 1,3-Propanediamine, N1,N1-dimethyl-N3-(5-methyl-5H-quindolin-11-yl)- (CA INDEX NAME)

Me2N- (CH2)3-NH

- RN 854190-76-4 CAPLUS
- CN 5H-Quindoline-11-carboxamide, N,N'-[1,3-propanediylbis[(methylimino)-2,1-ethanediyl]]bis[5-methyl- (9CI) (CA INDEX NAME)

PAGE 1-B

- RN 854190-81-1 CAPLUS
- CN 5H-Quindoline-11-carboxamide, N,N'-[1,2-ethanediylbis[(methylimino)-2,1ethanediyl]]bis[5-methyl- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 854190-92-4 CAPLUS

CN 1,3-Propanediamine, N,N'-dimethyl-N,N'-bis[2-[(5-methyl-5H-quindolin-11yl)amino]ethyl]- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

RN 854190-96-8 CAPLUS

 $\texttt{CN} \\ \hspace*{0.2cm} 1, 2-\texttt{Ethanediamine}, \hspace*{0.2cm} \texttt{N,N'-dimethyl-N,N'-bis} \\ [2-[(5-\texttt{methyl-5H-quindolin-11-methyl-N,N'-bis}]] \\ \hspace*{0.2cm} [2-[(5-\texttt{met$ 

yl)amino]ethyl]- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

ΝH

- IT 854190-49-1 854190-79-7 854190-83-3
  854190-94-6 854190-98-0
  RL: RCT (Reactant); RACT (Reactant or reagent)
  (preparation of antitumor quinoline derivs.)
- RN 854190-49-1 CAPLUS
- CN 5H-Quindoline-11-carboxamide, N-[3-[(3-aminopropyl)methylamino]propyl]-5-methyl- (CA INDEX NAME)

- RN 854190-79-7 CAPLUS
- CN 5H-Quindoline-11-carboxamide, N-[2-[[3-[(2aminoethy)]methylamino]propyl]methylamino]ethyl]-5-methyl-NAME)

- RN 854190-83-3 CAPLUS
- CN 5H-Quindoline-11-carboxamide, N-[2-[[2-[(2aminoethyl)methylamino]ethyl]methylamino]ethyl]-5-methyl- (CA INDEX NAME)

- RN 854190-94-6 CAPLUS
- CN 1,3-Propanediamine, N1-(2-aminoethyl)-N1,N3-dimethyl-N3-[2-[(5-methyl-5H-quindolin-11-yl)amino]ethyl]- (CA INDEX NAME)

H2N-CH2-CH2-N-(CH2)3-N-CH2-CH2-NH

- RN 854190-98-0 CAPLUS
- CN 1,2-Ethanediamine, N1-(2-aminoethy1)-N1,N2-dimethy1-N2-[2-[(5-methy1-5H-quindolin-11-y1)amino]ethy1]- (CA INDEX NAME)

H2N-CH2-CH2-N-CH2-CH2-N-CH2-CH2-NH

- II 854190-58-2P 854190-60-6P RL: RCI (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
  - (preparation of antitumor quinoline derivs.)
- RN 854190-58-2 CAPLUS
- CN Carbamic acid, [3-[methyl[3-[(5-methyl-5H-quindolin-11-yl) amino]propyl]amino[propyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

- RN 854190-60-6 CAPLUS
- CN 1,3-Propanediamine, N1-(3-aminopropy1)-N1-methy1-N3-(5-methy1-5H-quindolin-11-y1)- (CA INDEX NAME)

REFERENCE COUNT: 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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(FILE 'HOME' ENTERED AT 11:14:59 ON 26 JUN 2009)

FILE 'REGISTRY' ENTERED AT 11:15:33 ON 26 JUN 2009

L1 STRUCTURE UPLOADED

L2 1 S L1 L3 37 S L1 FULL

STR

FILE 'CAPLUS' ENTERED AT 11:16:11 ON 26 JUN 2009 L4 2 S L3

=> d 11 L1 HAS NO ANSWERS L1

G1 O, N

Structure attributes must be viewed using STN Express query preparation.